Confirmation No.: 2640

IN THE CLAIMS:

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1. (Presently amended) A <u>computer-implemented</u> method for calculating a global hydrophobic moment of a tertiary protein structure comprising a plurality of residues, the <u>method</u> comprising executing, via a computer, the following steps of:

calculating a centroid of residue centroids;

using the centroid of residue centroids as a spatial origin of a global linear hydrophobic moment;

calculating a first-order hydrophobic moment;

enhancing correlation between residue centroid magnitude and residue solvent accessibility, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using a distance metric;

using the first-order hydrophobic moment and the enhanced correlation between residue centroid magnitude and residue solvent accessibility to define the global linear hydrophobic moment, wherein each of the residue centroids contributes a magnitude and direction to the global linear hydrophobic moment, and wherein each residue centroid having a same fractional distance to a surface of the tertiary protein structure contributes an equivalent magnitude to the global linear hydrophobic moment by mapping each residue at a same distance from a center of the protein structure;

using the global linear hydrophobic moment to characterize an amphiphilicity of a tertiary protein structure; and

outputting the global linear hydrophobic moment to a user.

2. (Canceled)

- 3. (Original) The method of claim 1, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using an ellipsoidal metric.
- 4. (Original) The method of claim 1, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using a solvent accessibility metric.

Confirmation No.: 2640

5. (Original) The method of claim 1, wherein the centroid of residue centroids represents a geometric center of the tertiary protein structure.

- 5 6. (Cancelled)
 - 7. (Original) The method of claim 1, wherein the global linear hydrophobic moment characterizes a magnitude of amphiphilicity of the tertiary protein structure.
- 8. (Original) The method of claim 1, wherein the global linear hydrophobic moment characterizes a direction of amphiphilicity of the tertiary protein structure.
 - 9. (Original) The method of claim 1, wherein the global linear hydrophobic moment is used to identify functional regions of the tertiary protein structure.
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- 10. (Cancelled)
- 11. (Cancelled)
- 20 12. (Cancelled)
 - 13. (Cancelled)
- 14. (Presently amended) An apparatus for calculating a global hydrophobic moment 25 of a tertiary protein structure comprising a plurality of residues, the apparatus comprising:

a memory; and

at least one processor operative to:

calculate a centroid of residue centroids;

use the centroid of residue centroids as a spatial origin of a global linear hydrophobic moment;

Confirmation No.: 2640

calculate a first-order hydrophobic moment;

enhance correlation between residue centroid magnitude and residue solvent accessibility, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using a distance metric;

use the first-order hydrophobic moment and the enhanced correlation between residue centroid magnitude and residue solvent accessibility to define the global linear hydrophobic moment, wherein each of the residue centroids contributes a magnitude and direction to the global linear hydrophobic moment, and wherein each residue centroid having a same fractional distance to a surface of the tertiary protein structure contributes an equivalent magnitude to the global linear hydrophobic moment by mapping each residue at a same distance from a center of the protein structure;

use the global linear hydrophobic moment to characterize an amphiphilicity of a tertiary protein structure; and

output the global linear hydrophobic moment to a user.

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- 15. (Original) The apparatus of claim 14, wherein the centroid of the residue centroids represents a geometric center of the tertiary protein structure.
- 16. (Cancelled)

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- 17. (Original) The apparatus of claim 14, wherein the global linear hydrophobic moment is used to identify functional regions of the tertiary protein structure.
- 18. (Canceled)

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19. (Original) The apparatus of claim 14, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using an ellipsoidal metric.

Confirmation No.: 2640

20. (Original) The apparatus of claim 14, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using a solvent accessibility metric.

5 21. (Presently amended) An article of manufacture for calculating a global hydrophobic moment of a tertiary protein structure comprising a plurality of residues, comprising:

a computer-readable medium having computer-readable code embodied thereon, the computer-readable code comprising:

a step to calculate a centroid of residue centroids;

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a step to use the centroid of residue centroids as a spatial origin of a global linear hydrophobic moment;

a step to calculate a first-order hydrophobic moment;

a step to enhance correlation between residue centroid magnitude and residue solvent accessibility, wherein the correlation between residue centroid magnitude and residue solvent accessibility is enhanced using a distance metric;

a step to use the first-order hydrophobic moment and the enhanced correlation between residue centroid magnitude and residue solvent accessibility to define the global linear hydrophobic moment, wherein each of the residue centroids contributes a magnitude and direction to the global linear hydrophobic moment, and wherein each residue centroid having a same fractional distance to a surface of the tertiary protein structure contributes an equivalent magnitude to the global linear hydrophobic moment by mapping each residue at a same distance from a center of the protein structure;

a step to use the global linear hydrophobic moment to characterize an amphiphilicity of a tertiary protein structure; and

a step to output the global linear hydrophobic moment to a user.